

Amendments to the Claims

This listing of claims will replace all prior versions and listings of claims in the application:

Listing of Claims:

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1. (Withdrawn) A method for treatment of chronic pain comprising orally administering the composition of claim 9.
2. (Withdrawn) The method of claim 1 wherein said tricyclic antidepressant is administered in a dosage of from about 2.5 mg to about 25 mg daily.
3. (Withdrawn) The method of claim 2 wherein said tricyclic antidepressant compound is selected from the group consisting of doxepin, amitriptyline, desipramine, imipramine and physiologically acceptable acid addition salts thereof.
4. (Withdrawn) The method of claim wherein said physiologically acceptable acid addition salts are selected from the group consisting of the hydrochloride, hydrobromide, hydroiodide, acetate, valerate and oleate.
5. (Withdrawn) The method of claim 1 wherein said non-narcotic analgesic is administered in a dosage from about 0.50 gms to about 2.6 gms daily.
7. (Withdrawn) The method of claim 2 wherein said low dose of tricyclic antidepressant compound and said standard dose of non-narcotic analgesic are present in a single composition including a pharmaceutically acceptable vehicle for oral administration.
8. (Withdrawn) The method of claim 7 wherein said composition is in a form selected from the group consisting of tablets, capsules, caplets, oral solutions, and oral suspensions.
9. (Previously presented) A composition for treatment of chronic pain consisting essentially of a combination of a low dose of a tricyclic antidepressant compound and a standard dose of a non-narcotic analgesic in a pharmaceutical acceptable vehicle for oral administration.
10. (Previously presented) The composition of claim 9 being provided in a daily dosage form wherein said tricyclic antidepressant compound is present in an amount of about 2.5 mg to 25 mg.
11. (Original) The composition of claim 9 wherein said tricyclic antidepressant compound is selected from the group consisting of doxepin, amitriptyline, desipramine, imipramine, and physiologically acceptable acid addition salts thereof.

12. (Original) The composition of claim 9 wherein said physiologically acceptable acid addition salts are selected from the group consisting of the hydrochloride, hydrobromide, hydroiodide, acetate, valerate and oleate.

13. (Previously presented) The composition of claim 9 wherein said non-narcotic analgesic is administered in a dosage of from about 0.50 gms to about 2.6 gms daily.

14. (Previously presented) The composition of claim 9 wherein said non-narcotic analgesic is selected from the group consisting of acetaminophen or non-steroidal anti-inflammatory drugs.

15. (Original) The composition of claim 7 wherein the combination of a tricyclic antidepressant and a non-narcotic analgesic and a pharmaceutically acceptable vehicle is in a form selected from the group consisting of tablets, capsules, caplets, oral solutions and oral suspensions.